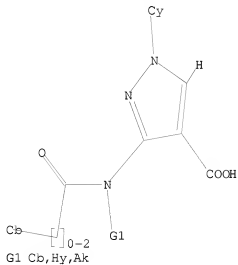


L1           STRUCTURE UPLOADED

=> d

L1 HAS NO ANSWERS

L1           STR



Structure attributes must be viewed using STN Express query preparation.

=> s l1 full

FULL SEARCH INITIATED 14:58:35 FILE 'REGISTRY'

FULL SCREEN SEARCH COMPLETED -       520 TO ITERATE

100.0% PROCESSED           520 ITERATIONS

331 ANSWERS

SEARCH TIME: 00.00.01

L2           331 SEA SSS FUL L1

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L2   ANSWER 1 OF 331   REGISTRY   COPYRIGHT 2009 ACS on STN

RN   1026248-17-8   REGISTRY

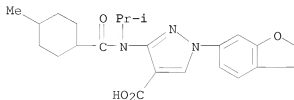
ED   Entered STN:   08 Jun 2008

CN   INDEX NAME NOT YET ASSIGNED

MF   C23 H29 N3 O4

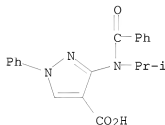
SR   Other Sources

Database: ChemSpider (ChemZoo, Inc.)



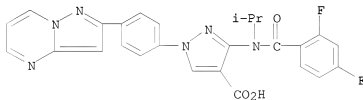
\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

L2 ANSWER 2 OF 331 REGISTRY COPYRIGHT 2009 ACS on STN  
RN 1026009-81-3 REGISTRY  
ED Entered STN: 06 Jun 2008  
CN 1H-Pyrazole-4-carboxylic acid, 3-[benzoyl(1-methylethyl)amino]-1-phenyl-  
(CA INDEX NAME)  
MF C20 H19 N3 O3  
SR Other Sources  
Database: ChemSpider (ChemZoo, Inc.)



\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

L2 ANSWER 3 OF 331 REGISTRY COPYRIGHT 2009 ACS on STN  
RN 934333-11-6 REGISTRY  
ED Entered STN: 06 May 2007  
CN 1H-Pyrazole-4-carboxylic acid, 3-[(2,4-difluorobenzoyl)(1-methylethyl)amino]-1-[4-(pyrazolo[1,5-a]pyrimidin-2-yl)phenyl]- (CA INDEX NAME)  
OTHER NAMES:  
CN 3-[(2,4-Difluorophenyl)carbonyl](1-methylethyl)amino]-1-[4-(pyrazolo[1,5-a]pyrimidin-2-yl)phenyl]-1H-pyrazole-4-carboxylic acid  
MF C26 H20 F2 N6 O3  
SR CA  
LC STN Files: CA, CAPLUS



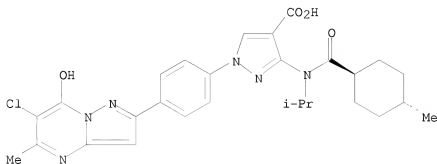
\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

1 REFERENCES IN FILE CA (1907 TO DATE)  
1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L2 ANSWER 4 OF 331 REGISTRY COPYRIGHT 2009 ACS on STN

RN 934333-10-5 REGISTRY  
 ED Entered STN: 06 May 2007  
 CN 1H-Pyrazole-4-carboxylic acid, 1-[4-(6-chloro-7-hydroxy-5-methylpyrazolo[1,5-a]pyrimidin-2-yl)phenyl]-3-[[1-(trans-4-methylcyclohexyl)carbonyl](1-methylethyl)amino]- (CA INDEX NAME)  
 FS STEREOSEARCH  
 MF C28 H31 Cl N6 O4  
 SR CA  
 LC STN Files: CA, CAPLUS

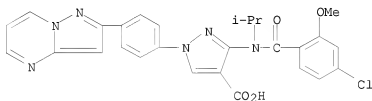
Relative stereochemistry.



\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

1 REFERENCES IN FILE CA (1907 TO DATE)  
 1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L2 ANSWER 5 OF 331 REGISTRY COPYRIGHT 2009 ACS on STN  
 RN 934333-09-2 REGISTRY  
 ED Entered STN: 06 May 2007  
 CN 1H-Pyrazole-4-carboxylic acid, 3-[[4-chloro-2-methoxybenzoyl](1-methylethyl)amino]-1-(4-pyrazolo[1,5-a]pyrimidin-2-ylphenyl)- (CA INDEX NAME)  
 OTHER NAMES:  
 CN 3-[[[4-Chloro-2-(methoxy)phenyl]carbonyl](1-methylethyl)amino]-1-[4-(pyrazolo[1,5-a]pyrimidin-2-yl)phenyl]-1H-pyrazole-4-carboxylic acid  
 MF C27 H23 Cl N6 O4  
 SR CA  
 LC STN Files: CA, CAPLUS

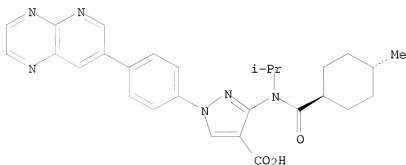


\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

1 REFERENCES IN FILE CA (1907 TO DATE)  
 1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L2 ANSWER 6 OF 331 REGISTRY COPYRIGHT 2009 ACS on STN  
 RN 934333-08-1 REGISTRY  
 ED Entered STN: 06 May 2007  
 CN 1H-Pyrazole-4-carboxylic acid, 3-[[{(trans-4-methylcyclohexyl)carbonyl}(1-methylethyl)amino]-1-(4-pyrido[2,3-b]pyrazin-7-yl)phenyl]- (CA INDEX NAME)  
 OTHER NAMES:  
 CN 3-[[{(trans-4-Methylcyclohexyl)carbonyl}(1-methylethyl)amino]-1-[4-(pyrido[2,3-b]pyrazin-7-yl)phenyl]-1H-pyrazole-4-carboxylic acid  
 FS STEREOSEARCH  
 MF C28 H30 N6 O3  
 SR CA  
 LC STN Files: CA, CAPLUS

Relative stereochemistry.

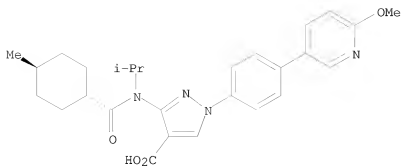


\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

1 REFERENCES IN FILE CA (1907 TO DATE)  
 1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L2 ANSWER 7 OF 331 REGISTRY COPYRIGHT 2009 ACS on STN  
 RN 934333-07-0 REGISTRY  
 ED Entered STN: 06 May 2007  
 CN 1H-Pyrazole-4-carboxylic acid, 1-[4-(6-methoxy-3-pyridinyl)phenyl]-3-[[{(trans-4-methylcyclohexyl)carbonyl}(1-methylethyl)amino]- (CA INDEX NAME)  
 OTHER NAMES:  
 CN 3-[[{(trans-4-Methylcyclohexyl)carbonyl}(1-methylethyl)amino]-1-[4-[6-(methyloxy)-3-pyridinyl]phenyl]-1H-pyrazole-4-carboxylic acid  
 FS STEREOSEARCH  
 MF C27 H32 N4 O4  
 SR CA  
 LC STN Files: CA, CAPLUS

Relative stereochemistry.

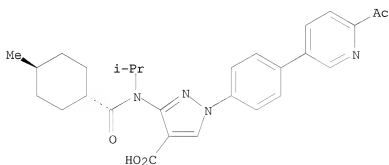


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1 REFERENCES IN FILE CA (1907 TO DATE)  
1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L2 ANSWER 8 OF 331 REGISTRY COPYRIGHT 2009 ACS on STN  
RN 934333-06-9 REGISTRY  
ED Entered STN: 06 May 2007  
CN 1H-Pyrazole-4-carboxylic acid, 1-[4-(6-acetyl-3-pyridinyl)phenyl]-3-  
[[4-(6-methylcyclohexyl)carbonyl](1-methylethyl)amino]- (CA INDEX  
NAME)  
OTHER NAMES:  
CN 1-[4-(6-Acetyl-3-pyridinyl)phenyl]-3-[[4-(6-methylcyclohexyl)carbonyl](1-methylethyl)amino]-1H-pyrazole-4-carboxylic  
acid  
FS STEREOSEARCH  
MF C28 H32 N4 O4  
SR CA  
LC STN Files: CA, CAPLUS

Relative stereochemistry.



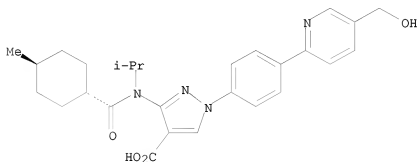
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1 REFERENCES IN FILE CA (1907 TO DATE)  
1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L2 ANSWER 9 OF 331 REGISTRY COPYRIGHT 2009 ACS on STN  
RN 934333-05-8 REGISTRY

ED Entered STN: 06 May 2007  
 CN 1H-Pyrazole-4-carboxylic acid, 1-[4-[5-(hydroxymethyl)-2-pyridinyl]phenyl]-  
 3-[[ (trans-4-methylcyclohexyl)carbonyl] (1-methylethyl)amino]- (CA INDEX  
 NAME)  
 OTHER NAMES:  
 CN 1-[4-[5-(Hydroxymethyl)-2-pyridinyl]phenyl]-3-[[ (trans-4-  
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 FS STEREOSEARCH  
 MF C27 H32 N4 O4  
 SR CA  
 LC STN Files: CA, CAPLUS

Relative stereochemistry.

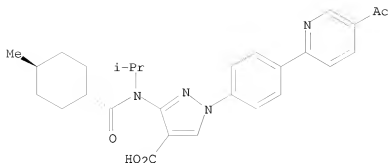


\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

1 REFERENCES IN FILE CA (1907 TO DATE)  
 1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L2 ANSWER 10 OF 331 REGISTRY COPYRIGHT 2009 ACS on STN  
 RN 934333-04-7 REGISTRY  
 ED Entered STN: 06 May 2007  
 CN 1H-Pyrazole-4-carboxylic acid, 1-[4-(5-acetyl-2-pyridinyl)phenyl]-3-  
 [[ (trans-4-methylcyclohexyl)carbonyl] (1-methylethyl)amino]- (CA INDEX  
 NAME)  
 OTHER NAMES:  
 CN 1-[4-(5-Acetyl-2-pyridinyl)phenyl]-3-[[ (trans-4-  
 methylcyclohexyl)carbonyl] (1-methylethyl)amino]-1H-pyrazole-4-carboxylic  
 acid  
 FS STEREOSEARCH  
 MF C28 H32 N4 O4  
 SR CA  
 LC STN Files: CA, CAPLUS

Relative stereochemistry.



\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

1 REFERENCES IN FILE CA (1907 TO DATE)  
1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

=> file caplus

COST IN U.S. DOLLARS

SINCE FILE

TOTAL

ENTRY

SESSION

FULL ESTIMATED COST

206.86

207.96

FILE 'CAPLUS' ENTERED AT 14:59:28 ON 17 MAR 2009

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FILE COVERS 1907 - 17 Mar 2009 VOL 150 ISS 12

FILE LAST UPDATED: 16 Mar 2009 (20090316/ED)

Caplus now includes complete International Patent Classification (IPC) reclassification data for the third quarter of 2008.

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This file contains CAS Registry Numbers for easy and accurate substance identification.

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L3 2 L2

=> d 13 1-2 ibib abs hitstr

L3 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2009 ACS on STN  
 ACCESSION NUMBER: 2007:409486 CAPLUS <<LOGINID:20090317>>  
 DOCUMENT NUMBER: 146:421977  
 TITLE: Preparation of 4-carboxypyrazoles as antivirals for treatment of hepatitis C virus (HCV) infection  
 INVENTOR(S): Bravi, Gianpaolo; Cheasty, Anne Goretti; Corfield, John Andrew; Grimes, Richard Martin; Harrison, David; Hartley, Charles David; Howes, Peter David; Medhurst, Katrina Jane; Meeson, Malcolm Lees; Mordaunt, Jacqueline Elizabeth; Shah, Pritom; Slater, Martin John; White, Gemma Victoria  
 PATENT ASSIGNEE(S): Smithkline Beecham Corporation, USA  
 SOURCE: PCT Int. Appl., 275pp.  
 CODEN: PIXXD2  
 DOCUMENT TYPE: Patent  
 LANGUAGE: English  
 FAMILY ACC. NUM. COUNT: 1  
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2007039146	A1	20070412	WO 2006-EP9238	20060921
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HN, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LT, LU, LV, LY, MA, MD, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RS, RU, SC, SD, SE, SG, SK, SL, SM, SV, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
PRIORITY APPLN. INFO.:			GB 2005-19474	A 20050923
			GB 2006-5467	A 20060317
OTHER SOURCE(S): MARPAT 146:421977				
GI				

\* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT \*

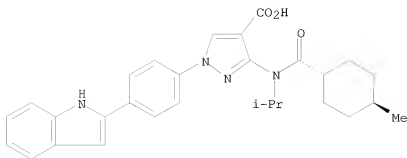
AB Title compds. [I; R1 = -Rx-Ry; Rx = (un)substituted phenylene, or 5- to 6-membered heteroarylene bonded through a ring C to the N of the pyrazole; Ry = heteroaryl, bonded such that when Rx = Ph, Ry is in the para-position; R2 = Ph, cycloalkyl, cycloalkylmethyl; R3 = heterocyclyl, (un)substituted Ph, alkyl; and their salts, solvates and esters], were prepared for treating hepatitis C virus (HCV) infection. Thus, a multi-step synthesis from Et 3-amino-1H-pyrazole-4-carboxylate via Et 3-[[[(trans-4-methylcyclohexyl)carbonyl](1-methylethyl)amino]-1H-pyrazole-4-carboxylate (preparation given) was provided for carboxypyrazole II. I inhibited HCV RNA-dependent RNA polymerase with IC50 <10 nM.

RN 934331-05-2 CAPLUS

CN 1H-Pyrazole-4-carboxylic acid, 1-[4-(1H-indol-2-yl)phenyl]-3-[[[(trans-4-methylcyclohexyl)carbonyl](1-methylethyl)amino]- (CA INDEX NAME)

Relative stereochemistry.

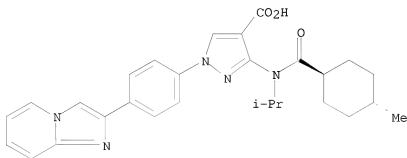




RN 934331-06-3 CAPLUS

CN 1H-Pyrazole-4-carboxylic acid, 1-(4-imidazo[1,2-a]pyridin-2-ylphenyl)-3-[[[(trans-4-methylcyclohexyl)carbonyl](1-methylethyl)amino]- (CA INDEX NAME)

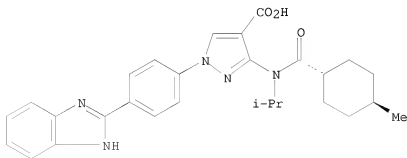
Relative stereochemistry.



RN 934331-07-4 CAPLUS

CN 1H-Pyrazole-4-carboxylic acid, 1-[4-(1H-benzimidazol-2-yl)phenyl]-3-[[[(trans-4-methylcyclohexyl)carbonyl](1-methylethyl)amino]- (CA INDEX NAME)

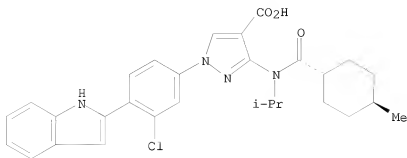
Relative stereochemistry.



RN 934331-08-5 CAPLUS

CN 1H-Pyrazole-4-carboxylic acid, 1-[3-chloro-4-(1H-indol-2-yl)phenyl]-3-[[[(trans-4-methylcyclohexyl)carbonyl](1-methylethyl)amino]- (CA INDEX NAME)

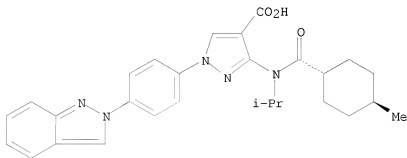
Relative stereochemistry.



RN 934331-09-6 CAPLUS

CN 1H-Pyrazole-4-carboxylic acid, 1-[4-(2H-indazol-2-yl)phenyl]-3-[[trans-4-methylcyclohexyl]carbonyl]-(1-methylethyl)amino]- (CA INDEX NAME)

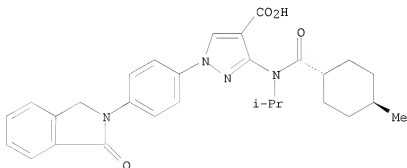
Relative stereochemistry.



RN 934331-10-9 CAPLUS

CN 1H-Pyrazole-4-carboxylic acid, 1-[4-(1,3-dihydro-1-oxo-2H-isoindol-2-yl)phenyl]-3-[[trans-4-methylcyclohexyl]carbonyl]-(1-methylethyl)amino]- (CA INDEX NAME)

Relative stereochemistry.

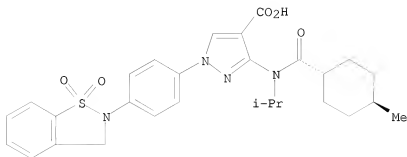


RN 934331-11-0 CAPLUS

CN 1H-Pyrazole-4-carboxylic acid, 1-[4-(1,1-dioxido-1,2-benzisothiazol-2(3H)-yl)phenyl]-3-[[trans-4-methylcyclohexyl]carbonyl]-(1-methylethyl)amino]-

(CA INDEX NAME)

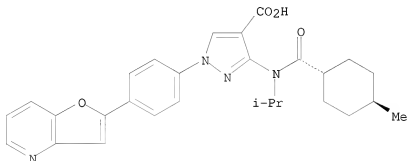
Relative stereochemistry.



RN 934331-12-1 CAPLUS

CN 1H-Pyrazole-4-carboxylic acid, 1-(4-furo[3,2-b]pyridin-2-ylphenyl)-3-[[4-(trans-4-methylcyclohexyl)carbonyl](1-methylethyl)amino]- (CA INDEX NAME)

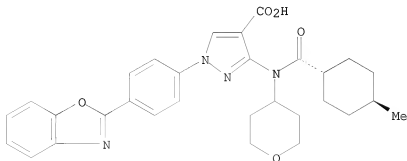
Relative stereochemistry.



RN 934331-13-2 CAPLUS

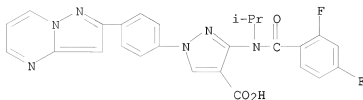
CN 1H-Pyrazole-4-carboxylic acid, 1-[4-(2-benzoxazolyl)phenyl]-3-[[4-(trans-4-methylcyclohexyl)carbonyl](tetrahydro-2H-pyran-4-yl)amino]- (CA INDEX NAME)

Relative stereochemistry.



RN 934333-11-6 CAPLUS

CN 1H-Pyrazole-4-carboxylic acid, 3-[(2,4-difluorobenzoyl)-1-methylethylamino]-1-(4-pyrazolo[1,5-a]pyrimidin-2-ylphenyl)- (CA INDEX NAME)



REFERENCE COUNT: 5 THERE ARE 5 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

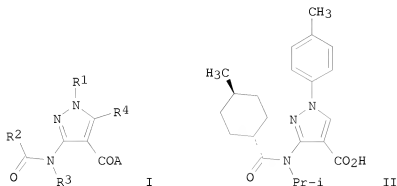
L3 ANSWER 2 OF 2 CAPLUS COPYRIGHT 2009 ACS on STN  
 ACCESSION NUMBER: 2005:10/5778 CAPLUS <<LOGINID::20090317>>  
 DOCUMENT NUMBER: 143:347168  
 TITLE: Preparation of pyrazole-4-carboxylic acids as HCV RNA polymerase inhibitors for the treatment of viral infections  
 INVENTOR(S): Bravi, Gianpaolo; Corfield, John Andrew; Grimes, Richard Martin; Guidetti, Rossella; Lovegrove, Victoria Lucy Helen; Mordaunt, Jacqueline Elizabeth; Shah, Pritom; Slater, Martin John  
 PATENT ASSIGNEE(S): Glaxo Group Limited, UK  
 SOURCE: PCT Int. Appl., 140 pp.  
 CODEN: PIXXD2  
 DOCUMENT TYPE: Patent  
 LANGUAGE: English  
 FAMILY ACC. NUM. COUNT: 1  
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2005092863	A1	20051006	WO 2005-GB1071	20050322
W:	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SM, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW			
RW:	BW, GH, GM, KE, LS, MM, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG			
EP 1730116	A1	20061213	EP 2005-729380	20050322
EP 1730116	B1	20070905		
R:	AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LI, LT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, HR, LV			
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AT 372323	T	20070915	AT 2005-729380	20050322
JP 2007530516	T	20071101	JP 2007-504463	20050322
ES 2292126	T3	20080301	ES 2005-729380	20050322
IN 2006DN05138	A	20070803	IN 2006-DN5138	20060906
US 20070167507	A1	20070719	US 2006-599346	20060926
PRIORITY APPLN. INFO.:			GB 2004-6916	A 20040326

GB 2004-6918	A	20040326
GB 2004-23000	A	20041015
GB 2004-23001	A	20041015
WO 2005-GB1071	W	20050322

OTHER SOURCE(S):  
GI

CASREACT 143:347168; MARPAT 143:347168



AB The invention relates to the title compds. I [wherein A = OH; R1 = (un)substituted (hetero)aryl or heterocyclyl; R2 = substituted Ph or (un)substituted cycloalkyl(methyl); R3 = (un)substituted heterocyclyl, heteroaryl, Ph or alkyl; R4 = H, and salts, solvates and esters thereof], a process for their prepns. and use as antiviral agents. For instance, acid II was synthesized by hydrolysis of the corresponding Et ester (preparation given) with LiOH in THF-EtOH at rt for 16 h. Exemplified compds. I (no indication) had inhibition against HCV RNA-dependent RNA polymerase with IC50 values of <80  $\mu$ M. Therefore, I and their pharmaceutical compns. are useful as HCV RNA polymerase inhibitors and in treating and/or preventing viral infections.

IT 865870-39-9P

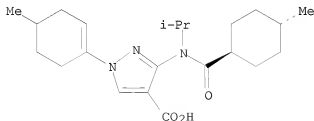
RL: PAC (Pharmacological activity); PUR (Purification or recovery); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(drug candidate; preparation of pyrazolecarboxylic acids as HCV RNA polymerase inhibitors for the treatment of viral infections)

RN 865870-39-9 CAPLUS

CN 1H-Pyrazole-4-carboxylic acid, 1-(4-methyl-1-cyclohexen-1-yl)-3-[[{(trans-4-methylcyclohexyl)carbonyl}(1-methylethyl)amino]- (CA INDEX NAME)

Relative stereochemistry.



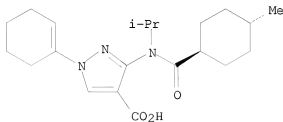
IT 865869-73-4P 865870-69-5P 865870-90-2P

RL: PAC (Pharmacological activity); RCT (Reactant); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); RACT (Reactant or reagent); USES (Uses)  
(drug candidate; preparation of pyrazolecarboxylic acids as HCV RNA polymerase inhibitors for the treatment of viral infections)

RN 865869-73-4 CAPLUS

CN 1H-Pyrazole-4-carboxylic acid, 1-(1-cyclohexen-1-yl)-3-[[1-(trans-4-methylcyclohexyl)carbonyl](1-methylethyl)amino]- (CA INDEX NAME)

Relative stereochemistry.

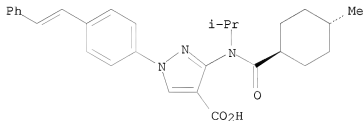


RN 865870-69-5 CAPLUS

CN 1H-Pyrazole-4-carboxylic acid, 3-[[1-(trans-4-methylcyclohexyl)carbonyl](1-methylethyl)amino]-1-[4-(2-phenylethenyl)phenyl]- (CA INDEX NAME)

Relative stereochemistry.

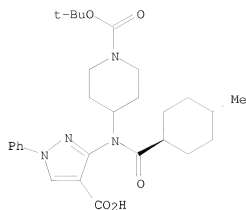
Double bond geometry unknown.



RN 865870-90-2 CAPLUS

CN 1-Piperidinecarboxylic acid, 4-[[4-(4-carboxy-1-phenyl-1H-pyrazol-3-yl)[1-(trans-4-methylcyclohexyl)carbonyl]amino]-, 1-(1,1-dimethylethyl) ester (CA INDEX NAME)

Relative stereochemistry.



IT 865869-71-2P 865869-72-3P 865869-74-5P  
 865869-75-6P 865869-76-7P 865869-77-8P  
 865869-78-9P 865869-79-0P 865869-80-3P  
 865869-81-4P 865869-82-5P 865869-83-6P  
 865869-84-7P 865869-85-8P 865869-86-9P  
 865869-87-0P 865869-88-1P 865869-89-2P  
 865869-90-5P 865869-91-6P 865869-92-7P  
 865869-93-8P 865869-94-9P 865869-95-0P  
 865869-96-1P 865869-97-2P 865869-98-3P  
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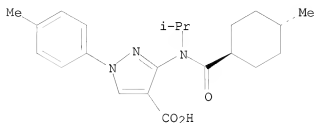
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(drug candidate; preparation of pyrazolecarboxylic acids as HCV RNA polymerase inhibitors for the treatment of viral infections)

RN 865869-71-2 CAPLUS

CN 1H-Pyrazole-4-carboxylic acid, 3-[[ (trans-4-methylcyclohexyl)carbonyl] (1-methylethyl)amino]-1-(4-methylphenyl)- (CA INDEX NAME)

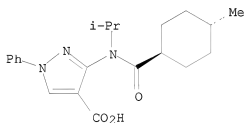
Relative stereochemistry.



RN 865869-72-3 CAPLUS

CN 1H-Pyrazole-4-carboxylic acid, 3-[[ (trans-4-methylcyclohexyl)carbonyl] (1-methylethyl)amino]-1-phenyl- (CA INDEX NAME)

Relative stereochemistry.



RN 865869-74-5 CAPLUS

CN 1H-Pyrazole-4-carboxylic acid, 1-(4-chloro-3-methylphenyl)-3-[[ (trans-4-methylcyclohexyl)carbonyl] (1-methylethyl)amino]- (CA INDEX NAME)

Relative stereochemistry.

